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                alerts (SDIs) affected
     10 DEC 17 COMPUAB reloaded; updating to resume; current-awareness
NEWS
                alerts (SDIs) affected
                SOLIDSTATE reloaded; updating to resume; current-awareness
NEWS
      11 DEC 17
                alerts (SDIs) affected
     12 DEC 17 CERAB reloaded; updating to resume; current-awareness
NEWS
                alerts (SDIs) affected
NEWS 13 DEC 17 THREE NEW FIELDS ADDED TO IFIPAT/IFIUDB/IFICDB
NEWS 14 DEC 30 EPFULL: New patent full text database to be available on STN
NEWS 15 DEC 30 CAPLUS - PATENT COVERAGE EXPANDED
     16 JAN 03
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                February 2005
NEWS EXPRESS JANUARY 10 CURRENT WINDOWS VERSION IS V7.01a, CURRENT
             MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
             AND CURRENT DISCOVER FILE IS DATED 10 JANUARY 2005
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0.21

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AND TECHNOLOGY CORPORATION, AND FACHINFORMATIONSZENTRUM KARLSRUHE

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SINCE FILE TOTAL
ENTRY SESSION

FULL ESTIMATED COST

0.06
0.27

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=> file re

'RE' IS AN AMBIGUOUS FILE OR CLUSTER NAME

REACTION - Reactions Cluster RESEARCH - Research Cluster

REGISTRY - The CAS Registry File of substances

ENTER FILE OR CLUSTER NAME (IGNORE): reg

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0.48

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STRUCTURE FILE UPDATES: 9 JAN 2005 HIGHEST RN 810659-29-1 DICTIONARY FILE UPDATES: 9 JAN 2005 HIGHEST RN 810659-29-1

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

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Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

=> s ENTER LOGIC EXPRESSION, QUERY NAME, OR (END):end SEARCH ENDED BY USER

```
=> e se-methyl-l-cysteine/cn
                   SE-METHYL P-METHOXYSELENOBENZOATE/CN
E1
                   SE-METHYL P-METHYLSELENOBENZOATE/CN
E2
             0 --> SE-METHYL-L-CYSTEINE/CN
E3
                   SE-METHYLSELENOCYSTEINE/CN
E4
                   SE-METHYLSELENOMETHIONINE/CN
E5
                   SE-N-DODECYL N, N-DIETHYLSELENOCARBAMATE/CN
E6
             1
                   SE-N-DODECYL O-ETHYL SELENOCARBONATE/CN
E7
             1
                   SE-N-HEXADECYL N, N-DIETHYLSELENOCARBAMATE/CN
E8
             1
                   SE-N-HEXADECYL O-ETHYL SELENOCARBONATE/CN
E9
             1
                   SE-N-OCTADECYL N, N-DIETHYLSELENOCARBAMATE/CN
E10
             1
                   SE-N-OCTADECYL O-ETHYL SELENOCARBONATE/CN
E11
             1
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E12

=> e4

L1

1 SE-METHYLSELENOCYSTEINE/CN

=> d 11

L1 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2005 ACS on STN

RN 26046-90-2 REGISTRY

CN L-Alanine, 3-(methylseleno)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Alanine, 3-(methylselenyl)-, L- (8CI)

OTHER NAMES:

CN 3-(Methylseleno)-L-alanine

CN Methylseleno-L-cysteine

CN Methylselenocysteine

CN Se-Methylselenocysteine

FS STEREOSEARCH

MF C4 H9 N O2 Se

LC STN Files: AGRICOLA, ANABSTR, BEILSTEIN\*, BIOSIS, CA, CAPLUS, CASREACT, CHEMCATS, CSCHEM, DDFU, DRUGU, TOXCENTER, USPATFULL (\*File contains numerically searchable property data)

DT.CA CAplus document type: Conference; Dissertation; Journal; Patent

RL.P Roles from patents: BIOL (Biological study); PREP (Preparation); PROC (Process); RACT (Reactant or reagent); USES (Uses)

RL.NP Roles from non-patents: ANST (Analytical study); BIOL (Biological study); FORM (Formation, nonpreparative); OCCU (Occurrence); PREP (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or reagent); USES (Uses)

Absolute stereochemistry.

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

99 REFERENCES IN FILE CA (1907 TO DATE)

99 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> file caplus

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 7.30 7.78

FULL ESTIMATED COST

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> l1 L299 L1 => 11/prep 99.L1 3245018 PREP/RL 10 L1/PREP L3 (L1 (L) PREP/RL) => save temp 12 secysts/a ANSWER SET L2 HAS BEEN SAVED AS 'SECYSTS/A' => d 13 5-10 ti fbib abs ANSWER 5 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN L3 A method of using synthetic L-Se-methylselenocysteine as a nutriceutical TI2002:364013 CAPLUS AN 136:369993 DN A method of using synthetic L-Se-methylselenocysteine as a nutriceutical TI Spallholz, Julian E.; Reid, Ted W.; Walkup, Robert D. IN Pharmase, Incorporated, USA PA Eur. Pat. Appl., 21 pp. SO

CODEN: EPXXDW

DT Patent

LA English

FAN.CNT 1

	PATENT NO EP 1205471				KIND DATE 20020515			APPLICATION NO.					DATE			
ΡI								EP 2001-103018				20010208				
	R:	AT,	BE,	CH,	DE,	DK, ES,	FR,	GB, GF	?, IT,	LI,	LU,	NL,	SE,	MC,	PT,	
		IE,	SI,	LT,	LV,	FI, RO,	MK,	CY, AI	TR							
								US	2000-	6775	63	7	A 2	0001	002	
	EP 1077209			A1	2001	0221	EP	2000-	1171	06		2	0000	809		
	R:	AT,	BE,	CH,	DE,	DK, ES,	FR,	GB; GF	R, IT,	LI,	LU,	NL,	SE,	MC,	PT,	
		IE,	SI,	LT,	LV,	FI, RO										
								US	1999-	3760	73	i	A 1	9990	816	
	US 2003083383			A1	2003	0501	US	2002-	2880	24		2	0021	105		
								US	1999-	3760	73	]	B2 1	.9990	816	
								US	2000-	6775	63	1	A3 2	0001	002	

OS CASREACT 136:369993

The invention describes the synthesis and use of L-Se-methylselenocysteine (I), a nutriceutical which is less toxic than L-selenomethionine towards normal cells. The synthesis involves mixing N-(tert-butoxycarbonyl)-L-serine with a dialkyl diazodicarboxylate and at least one of a trialkylphosphine, triarylphosphine and phosphite to form a mixture containing N-(tert-butoxycarbonyl)-L-serine  $\beta$ -lactone, addition of methylselenol or a salt, and deprotection. This synthesis significantly improves the manufacturing efficiency and utility I., a naturally occurring rare form of organic

selenium. I formed in this manner may be used as a nutriceutical in the diets of humans or animals for various beneficial purposes, such as, for example, to prevent or reduce the risk of developing cancer. A bar graph which compares the effect of I and L-selenomethionine on the growth of normal rabbit fibroblasts is given.

RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L3 ANSWER 6 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
- Synthesis of Novel Se-Substituted Selenocysteine Derivatives as Potential Kidney Selective Prodrugs of Biologically Active Selenol Compounds: Evaluation of Kinetics of  $\beta$ -Elimination Reactions in Rat Renal Cytosol
- AN 1996:241974 CAPLUS
- DN 124:306525
- Synthesis of Novel Se-Substituted Selenocysteine Derivatives as Potential Kidney Selective Prodrugs of Biologically Active Selenol Compounds: Evaluation of Kinetics of  $\beta$ -Elimination Reactions in Rat Renal Cytosol
- AU Andreadou, Ioanna; Menge, Wiro M. P. B.; Commandeur, Jan N. M.; Worthington, Eduard A.; Vermeulen, Nico P. E.
- CS Leiden Amsterdam Center for Drug Research, Vrije Universiteit Amsterdam, Amsterdam, 1081 HV, Neth.
- SO Journal of Medicinal Chemistry (1996), 39(10), 2040-6 CODEN: JMCMAR; ISSN: 0022-2623
- PB American Chemical Society
- DT Journal
- LA English
- Eighteen Se-substituted selenocysteine derivs. were prepared as potential AB kidney selective prodrugs which can be activated by renal cysteine conjugate β-lyase to selenium-containing chemoprotectants or antitumor agents. Selenocysteine derivs. with aliphatic and benzylic Se-substituents were synthesized by reducing selenocystine to selenocysteine followed by a reaction with the corresponding alkyl and benzyl halogenides. Selenocysteine derivs. with aromatic Se-substituents were synthesized by reaction of  $\beta$ -chloroalanine with substituted phenylselenol compds., which were formed by reducing substituted di-Ph diselenides by NaBH4. The enzyme kinetic parameters (apparent Km and Vmax) of the  $\beta$ -elimination reaction of the selenocysteine conjugates were studied in rat renal cytosol. The results suggest that Se-substituted L-selenocysteine conjugates are extremely good substrates for renal cysteine conjugate β-lyases as indicated by low apparent Km and high Vmax values. benzyl-substituted Se-conjugates appeared to be better substrates than the phenyl- and alkyl-substituted Se-conjugates. Corresponding L-cysteine S-conjugates were too poor substrates to obtain proper enzyme kinetics. Recently, local activation of cysteine S-conjugates by renal cysteine conjugate  $\beta$ -lyases was proposed as a new strategy to target antitumor agents to the kidney. Se-substituted selenocysteine conjugates may be more promising prodrugs because these are much better substrates for  $\beta$ -lyase.
- L3 ANSWER 7 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Preparation of sulfur and selenium amino acids with microbial pyridoxal phosphate enzymes
- AN 1988:128044 CAPLUS
- DN 108:128044
- TI Preparation of sulfur and selenium amino acids with microbial pyridoxal phosphate enzymes
- AU Esaki, Nobuyoshi; Soda, Kenji
- CS Inst. Chem. Res., Kyoto Univ., Uji, 611, Japan
- SO Methods in Enzymology (1987), 143 (Sulfur Sulfur Amino Acids), 291-7 CODEN: MENZAU; ISSN: 0076-6879
- DT Journal
- LA English
- AB The preparation of S-substituted L-homocysteines with L-methionine γ-lyase (I), S-substituted L-cysteines and Se-substituted L-selenocysteines with tryptophan synthase, L-selenocystine and -homocystine with O-acetylhomoserine sulfhydrylase, and deuterated and tritiated L-methionine and S-methyl-L-cysteine with I are illustrated.
- L3 ANSWER 8 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Selenium-containing amino acids
- AN 1984:4789 CAPLUS

100:4789 DN Selenium-containing amino acids TI Mitsui Toatsu Chemicals, Inc., Japan PA Jpn. Kokai Tokkyo Koho, 3 pp. SO CODEN: JKXXAF Patent DTJapanese LA FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE JP 58146286 A2 19830831 JP 1982-28108 19820225 ΡI JP 02054076 B4 19901120 JP 1982-28108 19820225 A composition containing methaneselenol [6486-05-1] or benzylselenol AB [16645-12-8] and L-serine [56-45-1] is treated with tryptophan synthetase [9014-52-2] to produce Se-methylselenocysteine [26046-90-2] or Sebenzylselenocysteine [2575-74-8]. Thus, a composition containing L-serine 30, methaneselenol 50, pyridoxal phosphate 0.01 mM, and tryptophan synthetase 10 mg/dL was shaken at 30° for 24 h. The medium contained Se-methylselenocysteine with a mol. yield rate of 28%. ANSWER 9 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN L3 Enzymatic synthesis of selenium-substituted L-selenocysteine with TI tryptophan synthase 1983:590469 CAPLUS AN 99:190469 DN Enzymatic synthesis of selenium-substituted L-selenocysteine with TI tryptophan synthase Esaki, Nobuyoshi; Tanaka, Hidehiko; Miles, Edith W.; Soda, Kenji AU Inst. Chem. Res., Kyoto Univ., Uji, 611, Japan CS FEBS Letters (1983), 161(2), 207-9 SO CODEN: FEBLAL; ISSN: 0014-5793 Journal DT English LA CASREACT 99:190469 OS When L-serine was incubated with the purified  $\alpha 2\beta 2$  complex of AB tryptophan synthase (EC 4.2.1.20) from Escherichia coli in the presence of a standard reaction mixture containing  $\alpha$ -tolueneselenol, Se-benzyl-L-5selenocysteine was formed with a yield of 44%, based on the L-serine used. The product was identified by several physicochem. criteria, including NMR. L-Serine was also converted to Se-methyl-L-selenocysteine by this method with methaneselenol as a reactant. The yield was 16%, based on L-serine. The reactivities of selenols were compared to those of thiols in a reaction system in which L-serine was used as a substrate. The specific activities of tryptophan synthase in β-replacement reactions with  $\alpha$ -tolueneselenol and methaneselenol were 0.96 and 0.77, resp., whereas those with  $\alpha$ -toluenethiol and methanethiol were 3.2 and 0.61, resp. Possible reasons for these reactivities are discussed. ANSWER 10 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN L3Selenoamino acids TI1979:522166 CAPLUS AN 91:122166 DN Selenoamino acids TISayuda, Kenji; Tanaka, Hidehiko IN Ajinomoto Co., Inc., Japan PA

SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION

JP 1977-117664 A 19770929 Eight selenoamino acids RSe(CH2)nCH(NH2)CO2H (R = organic residues; n = 1, 2) AB were prepared by reaction of R1(CH2)nCH(NH2)CO2H [R1 = halo, R2O (R2 = H, alkyl), R2S, R2SO, R2SO2] with RSeH in aqueous media in the presence of methioninase. Thus, Pseudomonas ovalis IFO 3738 was cultured on 1 kg of broth (pH 7.2) containing L-methionine 0.25, urea 0.1, peptone 0.1, glycerol 0.1, KH2PO4 0.1, K2HPO4 0.1, MgSO4.7H2O 0.01, and yeast extract 0.025 g/dL 18 h at 28° to give 2.2 kg cells, which were crushed in H3PO4 buffer and the supernatant treated on DEAE-cellulose and Sephadex G-200 to give 280 mg enzyme protein. A mixture of 0.1M L-methionine (in 0.2M H3PO4 buffer at pH 8.0), 0.1 mL 1M PhSeH (in EtOH), 0.5 mL 10-5M pyridoxal phosphate (in 0.02M H3PO4 buffer at pH 8.0), and 1 mL of the enzyme liquid (50 µg of protein/mL) was kept for 2 h at 37° under N with addition of 3 + 200  $\mu$ L of the enzyme liquid and 3 + 100  $\mu$ L the PhSeH liquid and the whole kept 25 min at 100° to give 4.3 mg  $\gamma$ -phenylseleno- $\alpha$ -aminobutyric acid [71128-79-5].

=> piperine

813 PIPERINE

9 PIPERINES

L4 815 PIPERINE

(PIPERINE OR PIPERINES)

=> d his

(FILE 'HOME' ENTERED AT 13:06:13 ON 11 JAN 2005)

FILE 'STNGUIDE' ENTERED AT 13:06:25 ON 11 JAN 2005

FILE 'HOME' ENTERED AT 13:06:29 ON 11 JAN 2005

FILE 'REGISTRY' ENTERED AT 13:06:46 ON 11 JAN 2005 E SE-METHYL-L-CYSTEINE/CN

L1 1 E4

FILE 'CAPLUS' ENTERED AT 13:08:00 ON 11 JAN 2005

L2 99 L1

L3 10 L1/PREP

SAVE TEMP L2 SECYSTS/A

L4 815 PIPERINE

=> 12 and 14

L5 1 L2 AND L4

=> d 15 ti fbib abs

L5 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN

TI Nutritional compositions containing selenium and lithium and use thereof as anti-HIV and anti-AIDS nutraceuticals and immunostimulants.

AN 2004:1074096 CAPLUS

DN 142:37306

TI Nutritional compositions containing selenium and lithium and use thereof as anti-HIV and anti-AIDS nutraceuticals and immunostimulants.

PA Serfontein, Willem Jacob, S. Afr.

SO PCT Int. Appl., 67 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 2004107881 A1 20041216 WO 2004-ZA60 20040603

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,

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LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
             NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
             TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
             AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
             EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
             SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
             SN, TD, TG
                                            ZA 2003-4360
                                                                 A 20030604
                                            ZA 2003-5112
                                                                 A 20030701
                                            ZA 2003-6713
                                                                 A 20030828
                                            ZA 2004-53
                                                                    20040106
     A nutrient composition or combination of compns. for the treatment or
AB
     prophylaxis of infections, in particular HIV/AIDS, and for the enhancement
     of immunity, based on selenium in synergistic combinations with biol.
     absorbable sources of glutathione, alkalinity enhancing components, a source of
     sulfur, an anti-mutagenic compound and for oral use, gastrointestinal
     absorption enhancers. Special uses relate to reducing risks of
     mother-to-child transmission and treating HIV-pos. pregnant women.
     Preferred further ingredients include antiinflammatory compds. and
     nutrients which control homocysteine.
              THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 10
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
=> black pepper
        232648 BLACK
          5541 BLACKS
        233735 BLACK
                 (BLACK OR BLACKS)
         10034 PEPPER
          2048 PEPPERS
         10731 PEPPER
                 (PEPPER OR PEPPERS)
           913 BLACK PEPPER
L6
                 (BLACK (W) PEPPER)
=> d his
     (FILE 'HOME' ENTERED AT 13:06:13 ON 11 JAN 2005)
     FILE 'STNGUIDE' ENTERED AT 13:06:25 ON 11 JAN 2005
     FILE 'HOME' ENTERED AT 13:06:29 ON 11 JAN 2005
     FILE 'REGISTRY' ENTERED AT 13:06:46 ON 11 JAN 2005
                E SE-METHYL-L-CYSTEINE/CN
L1
              1 E4
     FILE 'CAPLUS' ENTERED AT 13:08:00 ON 11 JAN 2005
             99 L1
L2
             10 L1/PREP
L3
                SAVE TEMP L2 SECYSTS/A
           815 PIPERINE
L4
              1 L2 AND L4
L5
          913 BLACK PEPPER
L6
=> 12 and 16
L7
        0 L2 AND L6
=> pepper
         10034 PEPPER
          2048 PEPPERS
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CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,

GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,

.W--

L8 10731 PEPPER

(PEPPER OR PEPPERS)

=> 12 and 18

L9 0 L2 AND L8

=> save temp all prosesht/l

L# LIST L1-L9 HAS BEEN SAVED AS 'PROSESHT/L'

=> logoff hold

COST IN U.S. DOLLARS

ENTRY
SESSION
FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

ENTRY
SESSION
ENTRY
SESSION
CA SUBSCRIBER PRICE

SINCE FILE
TOTAL
ENTRY
SESSION
-5.11

SESSION WILL BE HELD FOR 60 MINUTES
STN INTERNATIONAL SESSION SUSPENDED AT 13:29:03 ON 11 JAN 2005